## **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application.

## **Listing of Claims:**

- 1. (Currently Amended) A method for inhibiting platelet deposition in a patient in need thereof comprising administering a therapeutically effective amount of a nitric oxide adduct to the patient to inhibit platelet deposition; wherein the nitric oxide adduct is a nitrate—an angiotensin converting enzyme inhibitor which has at least one -O-NO<sub>2</sub> group, selected from the group consisting of a polypeptide; an amino acid; a sugar; an oligonucleotide; a branched or unbranched, saturated or unsaturated aliphatic hydrocarbon; an aromatic hydrocarbon; or a heterocyclic compound.
  - 2-4. (Cancelled)
- 5. (Original) The method of claim 1, further comprising administering at least one antithrombogenic compound or a therapeutic agent.
- 6. (Original) The method of claim 5, wherein the anti-thrombogenic compound is heparin, hirudin, an analog of hirudin, warfarin, aspirin, indomethacin, dipyridamole, prostacyclin, prostaglandin-E, a sulfinpyrazone, a phenothiazine, a RGD peptide, a RDG peptide mimetic, an agent that blocks platelet glycoprotein IIb-IIIa receptors, ticlopidine or clopidogrel.
- 7. (Original) The method of claim 5, wherein the therapeutic agent is a monoclonal antibody, a fragment of recombinant human protein, a viral vector or an anti-sense molecule.
- 8. (Currently Amended) A method for alleviating restenosis in a patient in need thereof comprising administering a therapeutically effective amount of a nitric oxide adduct to the patient to alleviate restenosis; wherein the nitric oxide adduct is a nitrate—an angiotensin converting enzyme inhibitor which has at least one -O-NO<sub>2</sub> group. selected from the group consisting of a polypeptide; an amino acid; a sugar; an oligonucleotide; a branched or unbranched, saturated or unsaturated aliphatic hydrocarbon; an aromatic hydrocarbon; or a heterocyclic compound.
  - 9-11. (Cancelled)
- 12. (Original) The method of claim 8, further comprising administering at least one anti-thrombogenic compound or a therapeutic agent.
- 13. (Original) The method of claim 12, wherein the anti-thrombogenic compound is heparin, hirudin, an analog of hirudin, warfarin, aspirin, indomethacin, dipyridamole,

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prostacyclin, prostaglandin-E, a sulfinpyrazone, a phenothiazine, a RGD peptide, a RDG peptide mimetic, an agent that blocks platelet glycoprotein IIb-IIIa receptors, ticlopidine or clopidogrel.

- 14. (Original) The method of claim 12, wherein the therapeutic agent is a monoclonal antibody, a fragment of recombinant human protein, a viral vector or an anti-sense molecule.
- 15. (Currently Amended) A method for treating a vascular complication caused by platelet deposition or thrombus formation in a patient in need thereof comprising administering a therapeutically effective amount of a nitric oxide adduct to the patient to treat the vascular complication caused by platelet deposition or thrombus formation; wherein the vascular complication caused by platelet deposition or thrombus formation is myocardial infarction, thrombophlebitis, thrombocytopenia or bleeding disorder; wherein the nitric oxide adduct is a nitrate—an angiotensin converting enzyme inhibitor which has at least one -O-NO<sub>2</sub> group. selected from the group consisting of a polypeptide; an amino acid; a sugar; an oligonucleotide; a branched or unbranched, saturated or unsaturated aliphatic hydrocarbon; an aromatic hydrocarbon; or a heterocyclic compound.
  - 16.-18. (Cancelled).
- 19. (Original) The method of claim 15, further comprising administering at least one anti-thrombogenic compound or a therapeutic agent.
- 20. (Original) The method of claim 19, wherein the anti-thrombogenic compound is heparin, hirudin, an analog of hirudin, warfarin, aspirin, indomethacin, dipyridamole, prostacyclin, prostaglandin-E, a sulfinpyrazone, a phenothiazine, a RGD peptide, a RDG peptide mimetic, an agent that blocks platelet glycoprotein IIb-IIIa receptors, ticlopidine or clopidogrel.
- 21. (Original) The method of claim 20, wherein the therapeutic agent is a monoclonal antibody, a fragment of recombinant human protein, a viral vector or an anti-sense molecule.
- 22. (Currently Amended) A method for inhibiting platelet deposition, platelet adhesion or thrombus formation in a patient in need thereof comprising administering a therapeutically effective amount of a nitric oxide adduct to the patient to treat a myocardial infarction, thrombophlebitis, thrombocytopenia or a bleeding disorder caused by the platelet deposition, platelet adhesion or the thrombus formation; wherein the nitric oxide adduct is a nitrate—an angiotensin converting enzyme inhibitor which has at least one -O-NO<sub>2</sub> group.

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selected from the group consisting of a polypeptide; an amino acid; a sugar; an oligonucleotide; a branched or unbranched, saturated or unsaturated aliphatic hydrocarbon; an aromatic hydrocarbon; or a heterocyclic compound.

23-25 (Cancelled)

- 26. (Original) The method of claim 22, further comprising administering at least one anti-thrombogenic compound or a therapeutic agent.
- 27. (Original) The method of claim 26, wherein the anti-thrombogenic compound is heparin, hirudin, an analog of hirudin, warfarin, aspirin, indomethacin, dipyridamole, prostacyclin, prostaglandin-E, a sulfinpyrazone, a phenothiazine, a RGD peptide, a RDG peptide mimetic, an agent that blocks platelet glycoprotein IIb-IIIa receptors, ticlopidine or clopidogrel.
- 28.(Original) The method of claim 26, wherein the therapeutic agent is a monoclonal antibody, a fragment of recombinant human protein, a viral vector or an anti-sense molecule.
- 29. (Currently Amended) A method for treating a dysfunction in the endothelium of a patient comprising administering a therapeutically effective amount of a nitric oxide adduct to the patient wherein the nitric oxide adduct is a nitrate—an angiotensin converting enzyme inhibitor which has at least one -O-NO<sub>2</sub> group, selected from the group consisting of a polypeptide; an amino acid; a sugar; an oligonucleotide; a branched or unbranched, saturated or unsaturated aliphatic hydrocarbon; an aromatic hydrocarbon; or a heterocyclic compound.

30-32 (Cancelled)

- 33. (Original) The method of claim 29, further comprising administering at least one anti-thrombogenic compound or a therapeutic agent.
- 34. (Original) The method of claim 33, wherein the anti-thrombogenic compound is heparin, hirudin, an analog of hirudin, warfarin, aspirin, indomethacin, dipyridamole, prostacyclin, prostaglandin-E, a sulfinpyrazone, a phenothiazine, a RGD peptide, a RDG peptide mimetic, an agent that blocks platelet glycoprotein IIb-IIIa receptors, ticlopidine or clopidogrel.
- 35. (Original) The method of claim 33, wherein the therapeutic agent is a monoclonal antibody, a fragment of recombinant human protein, a viral vector or an anti-sense molecule.